Construction of a Glucose Biosensor by Immobilizing Glucose Oxidase within a Poly(o-phenylenediamine) Covered Screen-printed Electrode

Miao Yuqing, Chen Jianrong, Wu Xiaohua
College of Chemistry and Life Science, Zhejiang Normal University
Jinhua 321004, China

Abstract: The glucose biosensors were prepared by the electropolymerization of the non-conductive polymer, Poly(o-phenylenediamine), onto a planar screen-printed electrode. A fabrication procedure was performed to decrease the waste of expensive enzyme. The amperometric glucose response was measured by the potentiostating of the prepared glucose biosensors at a potential of 0.3 V with ferrocene as mediator. Results show that the obtained biosensors have a linear range up to 25 mM glucose, fast response time (100s) and high sensitivities (16.6 nA/mM). Also, the effects of applied potential and sweeping number of Cyclic voltammograms for electropolymerization were systematically investigated and optimal values were recorded.

Key words: Poly(o-phenylenediamine), non-conducting polymer, biosensor, glucose

INTRODUCTION

Since the pioneering work of Clark and Lyons[1], the integration of an enzyme into an electrode attracted much interest in the development of biosensors up to now[2-6]. The advantages associated with these devices are safety, simplicity, minimal sample preparation, easy to handle, economy, accuracy, precision, and high sensitivity as well as the possibility of developing compact and portable analyzers[7,8].

In recent years, there has been a growing interest in immobilizing biomolecules in electropolymerized films to develop biosensors. The electropolymerization of electrically (non-)conducting polymers, such as poly(3-aminophenol), poly(1,3-diaminoarenzene), polyphenol, polyaniline, polyacletylne, polyindole or polypyrrole, is an interesting and effective procedure to prepare biosensors due to its simple preparation, easy miniaturization, and the precise localization of biomolecules[9-11]. Also the thickness of the resulting film could be controlled easily[12].

The electrochemical method involves the entrapment of biomolecules in organic polymers during their electrogeneration on an electrode surface. For most of research, the electrodes were dipped into the solution containing enzymes and monomer for copolymerization. One of the main disadvantages is that only a few enzymes in solution could be immobilized successfully and most expensive enzymes are disused[13,14].

In this research, a screen-printed planar electrode was used to fabricate the amperometric glucose biosensors by immobilizing GOD into the electropolymerized PPD film with ferrocene as the electron transfer mediator. Enzyme was dried on the surface of electrode before PPD film was formed. Therefore, most of enzyme was constrained under formed PPD film. In addition, only 20 μl enzyme was needed to cover the electrode surface. Few enzymes would be wasted in this study.

MATERIALS AND METHODS

Reagents: The enzyme glucose oxidase (GOD) from Aspergillus niger (EC 1.1.3.4, 158 units/mg of solid, type II) was obtained from Sigma Chemical Co. The monomer o-phenylenediamine (o-PD) for the electropolymerization was obtained from SCRC (Shanghai, China). PBS (pH 7.2) containing 0.01 M phosphate buffer, 0.137 M NaCl and 2.7 mM KCl was used as running buffer. GOD and ferrocene were prepared in PBS.

Apparatus: Electropolymerizations, amperometric measurements, and cyclic voltammetry (CV) were carried out with CHI 650A electrochemical workstation. Screen-printed electrodes were used for all electrochemical experiments. Working electrodes and counter electrodes were made of conductive carbon ink. One naked silver band was performed in 0.01 M FeCl₃ solution for 10 min to fabricate a reference electrode.

Preparation of glucose biosensors: Firstly, 20 μl 1mg/ml GOD and 20 μl 0.5 mM ferrocene were applied on the surface of electrodes. After drying, o-PD was dropped on the electrode to cover the below enzyme and ferrocene, and then the electropolymerization was performed by voltage cycling between –0.2 and 0.8 V vs Ag/AgCl and a scan rate of 50 mV/s for a certain number of cycles. Unless otherwise indicated, 20 μl of a 5 mM o-PD in PBS (pH 7.2) is used for the procedure. The resulting biosensors were rinsed with...
0.01 M PBS (pH 7.2) to remove un-immobilized enzymes, and were stored at 4 °C when not in use.

**Measurements:** Cyclic voltammogram measurements were carried out in 0.01 mM PBS (pH 7.2). Amperometric responses of the fabricated biosensors to glucose were studied by injections of aliquots of the glucose stock solution, and the resulting oxidation currents were monitored. All measurements were carried out at room temperatures.

**RESULTS and DISCUSSION**

The electropolymerization was performed by cyclic voltammetry between −0.2 and 0.8 V and a scan rate of 50 mV/s, using a 5 mM o-PD solution in 0.01 M PBS (pH 7.2). The results are shown in Fig. 1. The initial CV curve shows two irreversible oxidation peak at 0.35 V and 0.55 V corresponding to the polymerization of o-PD. On the successive cycles, however, the peak currents decreases, indicating that electropolymerization is a self-limiting process and the oxidation of o-PD may produce a compact and insulting film of PPD, which prevents from further deposition of polymer.[15] After first cycle, a pair of newly formed redox peaks could be observed at −0.2 and 0 V, which corresponds to the reduction and oxidation of PPD film. The detailed information of polymer structure and the polymerization mechanism could be found in the previous report.[16]

The enzyme should be kept in neutral pH for optimum activity, so a buffer of pH 7.2 was chose for the electropolymerization of PPD to cover the enzyme. Fig. 1b shows the CV of the electropolymerization of PPD with GOD presence on the electrode surface. Compared with Fig. 1a, two irreversible oxidation peaks at 0.35 V and 0.55 V shifts to more positive potential at 0.4 and 0.6 V in the presence of GOD. Also, the first oxidation peak decreases and the second oxidation peak increases due to the presence of GOD.

Fig. 2: Cyclic voltammograms of the PPD covered screen-printed electrodes with (a) only GOD; (b, c) GOD and ferrocene immobilized in PBS (pH 7.2) in the absence (a, b) and presence (c) of 35 mm glucose.

Fig. 2 shows the cyclic voltammograms obtained from the enzyme electrode in PBS (pH 7.2). Without glucose, the enzyme contributes no response and the GOD electrode only exhibits the quasi-reversible electrochemical behavior of ferrocene/ferricinium redox couple (Fig. 2b). The oxidation current is increased to a large extent in the presence of glucose (Fig. 2c), which is indicative of the enzyme-dependent catalytic reduction of the ferricinium ion produced at oxidizing potential[17]. Ferrocene molecules works well as an electron transfer mediator between the electrode substrate and the redox center of GOD.

The effect of applied potential on the biosensor response is shown in Fig. 3. The electrode response to glucose begins to increase at 0.1 V with the change of applied potential and becomes nearly constant at
potentials in the range of 0.3 to 0.6 V vs Ag/AgCl. The significant increased sensitivity with applied potential from 0.1 to 0.3 mV is attributed to the increased driving force for the oxidation of ferrocene to ferricinium. A further increase of the potential resulted in little change in current response. The potential of 0.3 V was selected for the following experiments so as to achieve the greatest sensitivity and relatively low background signal.

Fig. 4 Comparison of the current response of glucose biosensors with different concentration of ferrocene of (a) 0.1; (b) 0.25; (c) 0.5 and (d) 1 mM. Polarized potential, 0.3 V.

Normally, substrate (O\textsubscript{2}) or product (H\textsubscript{2}O\textsubscript{2}) of GOD enzyme reaction can be monitored for the development of glucose biosensors\textsuperscript{[18,19]}. The simplest method is to detect the consumption of oxygen at negative potentials (~0.6 V vs Ag/AgCl). However, the oxygen in the sample interferes the result of detection and such biosensors are not suitable for low analyte concentration. Another way is to detect of the oxidization of H\textsubscript{2}O\textsubscript{2}. A lot of reductants (for example, ascorbate, bilirubins, acetaminophen etc) in biological liquids are able to be oxidized at the same potential and produce high noise signal\textsuperscript{[20]}. Soluble or immobilized mediators have been developed to shuttle electron between active center of enzyme and electrode surface. Ferrocene was employed in this study for the development of glucose biosensor. Fig. 4 indicates that the current response of the biosensor increases sharply with the increase of the concentration of mediator from 0.1 to 0.5 mM. The biosensor response is limited by the enzyme-mediator kinetics at low mediator concentration and enzyme-substrate kinetics at high concentration of mediator. Higher levels of ferrocene concentration did not improve the signal response further and in fact caused the increase of background current, therefore, the concentration of ferrocene was set at 0.5 mM for all further work.

Fig. 5 The sensitivity of biosensors to glucose, which were prepared from (a) 2; (b) 3; (c) 4; (d) 5 circles of CV for PAN polymerization.

How the thickness of PPD film influences the final property of glucose biosensors was studied by changing the circle number of CV sweep for PPD polymerization (Fig. 5). Since most of charges for electropolymerization are consumed at the first several cycles, the 2, 3, 4, 5 cycles of CV were studied. When the scan cycle is only 2 cycles, the polymer film obtained is too loose or too thin to obtain a high amount of GOD. The biosensor prepared from 3 cycles of CV gives the highest sensitivity, and then the sensitivity decreases with the increase of cycles. It means that the non-conducting PPD film prevents the electron shuttle between GOD and electrode surface. Considering the sensitivity of biosensor, 4 circles of CV for PPD polymerization were chosen.

Fig. 6 The typical responses of a biosensor to glucose of 20 mM at 0.3 V.

When the glucose biosensor was polarized at 0.3 V vs Ag/AgCl and then a glucose solution was added into PBS (pH 7.2), a gradual increase in the oxidation current was observed (Fig. 6). The oxidation current
reached a stable steady-state value within 100 s. The repeatability of response of a typical glucose enzyme biosensor was investigated at a glucose concentration of 20 mM. The mean current was 231 nA with a rsd. of 6.2% (n=10). Fig. 7 shows the experimental calibration curve of the current response against glucose concentration with experimental condition chosen above. The linear range is 5-25 mM glucose and the sensitivity at 0.3 V is 16.6 nA/mM.

**CONCLUSION**

In this study, the electropolymerized non-conducting PPD film was used to immobilize GOD for the development of a glucose biosensor. GOD was dried on the surface of electrode before the electropolymerization of PPD. The procedure is simple and repeatable. Especially, the procedure decreases the waste of expensive enzyme.

**ACKNOWLEDGMENTS**

This material is based upon work funded by the National Natural Science Foundation of China (Grant No. 90406016), State Key Laboratory of Electroanalytical Chemistry of Chinese Academy of Sciences, Science and Technology Department of Zhejiang Provice.

**REFERENCES**